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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification : Not classified	A2	(11) International Publication Number: WO 99/00001 (43) International Publication Date: 7 January 1999 (07.01.99)
(21) International Application Number: PCT/KR98/00183 (22) International Filing Date: 26 June 1998 (26.06.98) (30) Priority Data: 1997/28473 27 June 1997 (27.06.97) KR 1997/28474 27 June 1997 (27.06.97) KR 1998/2226 24 January 1998 (24.01.98) KR (71) Applicant (for all designated States except US): LG CHEMICAL, LTD. [KR/KR]; LG Twin Towers, 20, Yoido-dong, Yongsungpo-gu, Seoul 150-721 (KR). (72) Inventors; and (75) Inventors/Applicants (for US only): AHN, Soon-Ho [KR/KR]; 107-1106, Lucky Hana Apt., Shinsung-dong, Yousung-gu, Taejon 305-345 (KR). CHUNG, Geun-Chang [KR/KR]; 6-103, LG Apt., Doryong-dong, Yousung-gu, Taejon 305-345 (KR). LIM, Mi-Ra [KR/KR]; 3-103, LG Chem. Dormitory, Doryong-dong, Yousung-gu, Taejon 305-340 (KR). JUN, Sung-Hui [KR/KR]; 3-103, LG Chem. Dormitory, Doryong-dong, Yousung-gu, Taejon 305-340 (KR). BAE, Joon-Sung [KR/KR]; 3-412, LG Chem. Dormitory, Doryong-dong, Yousung-gu, Taejon 305-340 (KR). KWAK, Mi-Seon [KR/KR]; 105-1110, Hanbat Garden, Sanjung-dong, Jung-gu, Taejon 301-211 (KR).		(74) Agent: KIM, Seong-Ki; Hubba Hubba Building, 7th floor, 648, Yoksam-dong, Kangnam-gu, Seoul 135-080 (KR). (81) Designated States: CN, JP, US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: LITHIUM ION SECONDARY BATTERY AND MANUFACTURING METHOD OF THE SAME (57) Abstract <p>The present invention discloses an improved lithium battery comprising a positive electrode, a negative electrode and a separator, and a manufacturing method of the same. The positive and negative electrodes comprise an active compounds selected from the group consisting of a compound being capable of reacting reversibly with a lithium ion, a compound having a structure that products of reacting a lithium ion with an electrolyte salt or an electrolyte solvent are capable of deposition or precipitation, or a compound having a structure that a lithium ion can be intercalated, an additive comprising a fiber of micron scale having an electron conductivity, a collector comprising metal or carbon material, and an adhesive. The lithium ion secondary battery prepared from the above described electrode has a low internal resistance, a high capacity and a high-speed recharging-discharging capability.</p>		

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LITHIUM ION SECONDARY BATTERY AND MANUFACTURING METHOD OF THE SAME

BACKGROUND OF THE INVENTION

Field of the Invention

5 The present invention relates to a lithium ion secondary battery having a low internal resistance and a greatly increased capacity, and more particularly to an improved lithium ion secondary battery constructed by selecting a composite electrode which is prepared by adding a conductive filament to electrode active materials during preparation of the battery, and a manufacturing method of the same.

10 Description of the Related Art

 While a lithium ion battery developed heretofore has a high energy density and a specific energy, it has a problem of high internal resistance. Accordingly, a practical capacity of the battery is prone to decline due to a rapid drop of battery voltage during discharging at a high speed. Because of such a disadvantage, it is known that lithium
15 battery is inefficient in electronic devices which require high current. One of the reasons of a high internal resistance and a slow recharging or discharging speed of the lithium battery is a low electrical conductivity of materials and structures of conventional electrodes.

 In general, an electrode of a lithium ion battery comprises powder type active
20 materials into which lithium ion can be intercalated and an adhesive polymer to fix the active material on a collector. Active materials such as carbonaceous materials as well as lithium metal oxides such LiCoO_2 , LiMn_2O_4 , LiNiO_2 have a poor electronic conductivity.

 Also, particles of these active materials expand and contract repeatedly when lithium ions are intercalated or deintercalated. Stress resulting from intercalation or
25 deintercalation along with swelling of an adhesive by an electrolyte causes loss of point contacts between particles of active materials. Accordingly, it is necessary to form electron delivery passages from collector to active materials while preparing the electrode for a high conductivity. In the case that the active materials of the anode are graphite or other type of carbon materials, the problem of conductivity is not as severe as in case of
30 cathode. However, an additional conductor needs to be added depending on the kinds of the battery and the objects for designing battery.

For this purpose, powders such as graphite in an average diameter of about 1 ~ 20 μ m or carbon black of an average size of about 10 ~ 500 nm can be added, as a conductor to the electrode. In these electrodes the conduction network is formed by the carbon powders residing in the void of active powder beds and interconnected by particle to particle point contacts. In order for the particles to function as a conductive matrix of electrode, contacts of active material particles should not be disconnected between collector and the forefront particles of electrode. However, these carbon materials have a lower electrical conductivity of about one thousandth compared to that of metal. And, since carbon particles in the electrode are electrically connected through multitude of point contacts, internal resistance within the electrode is fairly high.

The process for forming electron passages by a particle type conductor depends solely on the probability of encounter among particles. For the making of electron passages by addition of the powder to be successful, the probability of encounter must be high enough to reach a significant level. Also, conductivity through electron passages by point contacts of particles is proportional to the increase of contacting frequency. A good conductivity can not be obtained without forming multiplicity of electron passages.

In order to achieve these objectives under the prior art, a significant volumetric content of conductive additive has been used. Simple geometric calculations and common practice in Li-ion battery industry indicates the volume fraction of conductive carbon should range somewhere between about 6 to about 15 volume percent of the total electrode volume. Furthermore, the conductive particles must be fine enough to fit into the openings of active particle layer, resulting in the use of carbons with a small particle size and a large specific surface area. Naturally the electrode active surface area escalates unnecessarily, resulting in overall increase in reaction with electrolytes and organic solvents. The increased reactions can manifest as an irreversible loss of anode capacity, gradual oxidation and consumption of electrolyte on the cathode, which cause capacity decline during cycling, and an increased threat to battery safety by gas evolution and exothermic solvent oxidation.

In view of manufacturing the electrodes, the fine carbon particulates cause poor dispersion, which makes it difficult to get uniformly dispersed slurry. Further, the excess of conductors which do not contribute in forming electron passages

remains inside the electrode and constitutes an inactive volume, and thus limits the capacity of the battery and the energy density.

SUMMARY OF THE INVENTION

It is therefore an object of the invention to provide a rechargeable lithium ion battery comprising an electrode made of a fibriform electrical conductor of micron scale, and a manufacturing method of the same.

One strand of fiber made of active material particles has the same effects as those particles in a row. The surface area per unit volume of fibers is small, and the electric conductivity toward horizontal direction in fiber is superior to that of powders. Further, the mechanical strength of fiber is extraordinarily superior to that of the particles in a row.

The present invention provides a battery comprising an electrode having conductive electron passages embedded in it, wherein the electron passages are made of fibrous materials, the conductivity of the electrode is increased greatly.

BRIEF DESCRIPTION OF DRAWING

The present invention will be described with reference to the accompanying drawings of which:

Figure 1 shows scanning electron micrographs of metal fiber-containing electrodes, in which (A) is of LiCoO_2 cathode with 4 wt% of stainless steel 316L fiber; (B) is of LiCoO_2 cathode with 7 wt% of fiber; (C) and (D) are of MCMB anode with 0.5% of 316L fiber. The metal fibers are about 2 μm in diameter.

Figure 2 shows an influence of metal fiber addition on cathode discharge profile. The cathodes are composed of LiCoO_2 particles, PVdF binder, graphite conductive additive, and indicated weight percentage of metal fiber. Electrodes were discharged at 1/40 C rate. Voltage limit: 3.0 to 4.2 Volts vs. Li/Li^+ .

Figure 3 shows an impact of conductive additives on anode reactions with lithium ion. The volume fraction of conductive additives were maintained at approximately 0.8%. The electrodes were cycled at 0.1C rate between 0.01 and 1.5 V(vs. Li/Li^+). At least three electrode specimens were tested and the reversible and irreversible capacities calculated and appeared in the Figure. Cr and Cirr are defined as reversible and irreversible lithium ion uptake, respectively.

Figure 4 shows battery capacity behaviors with cycling. The particular cell with metal fiber, shown here, was fabricated using 316L fiber-containing anode(2% w/w) and regular cathode.

DETAILED DESCRIPTION OF THE INVENTION

5 The battery prepared by the method of the present invention has a low electrical resistance and a high-rate capability.

In order to achieve the objectives of the present invention, the lithium secondary battery provided according to the present invention comprises a cathode electrode, an anode electrode, an electrolyte and a separator.

10 In the battery of the present invention, the cathode electrode comprises: an active material which is capable of reacting reversibly with lithium ion; a compound having a structure in which reaction products of lithium ion with an electrolyte salt or an electrolyte solvent can be deposited or precipitated; or a compound having a structure with which lithium ion can be intercalated. Examples of such active materials include a
15 compound represented by: $\text{Li}_{1-x}\text{A}_x\text{Ni}_{1-y}\text{B}_y\text{O}_2$ (where, A = alkaline metal or alkaline earth metal, B = transition metal, $0 \leq x \leq 0.1$, $0 \leq y \leq 1.0$); $\text{LiMn}_{2-y}\text{M}_y\text{O}_4$ (where, M= Fe, Co, Ni: $0.02 \leq y \leq 0.3$); $\text{Li}_{1-x}\text{Ni}_{1-y}\text{B}_y\text{O}_2$ (where, B= transition metal, $0 \leq x \leq 0.1$, $0 \leq y \leq 1.0$); NbSe_3 ; $\text{Li}_x\text{V}_2\text{O}_5$; or $\text{Li}_x\text{V}_6\text{O}_{13}$ (where, $0 \leq x \leq 3.0$).

Also, the cathode electrode comprises an additive comprising an electron
20 conductive fiber of micron scale, a collector comprising metal and carbon materials, and an adhesive fixing the active materials to the collector.

The anode electrode comprises a material, as an active material, which is capable of reacting reversibly with lithium ion; a compound having a structure in which reaction products of lithium ion with an electrolyte salt or an electrolyte solvent can be deposited
25 or precipitated; or a compound having a structure with which lithium ion can be intercalated. The anode also comprises an additive comprising an electron conductive fiber of micron scale, an adhesive comprising polymer material, and a collector consisting of metal and carbon materials.

Among the active materials for the anode electrode, compounds having a structure
30 with which a lithium ion can be intercalated is classified into carbon group compounds and non-carbon group compounds. The carbon group compounds

are further classified into graphite group compounds and non-graphite group compounds. Examples of the graphite group compounds include flake type natural or synthesized graphite and a graphite based on mesophase pitch such as MCF, MCMB and the like. Examples of non-graphite group compounds include an amorphous carbon group
5 material thermally treated at 800~1800 °C, such as petroleum coke, soft carbon, hard carbon, polyparaphenylene or polyacenic compounds, and compounds based on coal. Examples of non-carbon group materials include metal oxides, metal sulfides, and metal selenides, also called dichalcogenides. Examples of metal oxides include SnO₂, WO₂, MoO₂ and the like. Examples of metal sulfides include TiS₂, MoS₂ and the like.
10 Examples of metal selenides include NbSe₃ and the like. While one or more active materials can be used, the kind of active materials is not limited in the present invention.

Conductive fibrous additive to the cathode and anode electrodes includes metal fiber and carbon fiber. Examples of metal fiber include iron, nickel, copper, zinc, titanium, aluminum, silver, gold, platinum, alloys such as iron-chromium alloy, iron-
15 chromium-nickel alloy (popular name: stainless steel), and aluminum alloy such as aluminum-copper, aluminum-manganese, aluminum-magnesium, aluminum-silicon-magnesium. Examples of carbon fiber include fibers based on synthesized graphite, active carbon or vapor-grown whisker. The diameter and the aspect ratio of preferable fibriform depend on the particle size distribution of the electrode active materials. In
20 general, a fiber with diameter from about 0.1 microns to about 25 microns and the aspect ratio of about 4 to about 2500 is preferable.

These fibers could be used without further treatment, or with surface treatment to increase adhesion strength. Examples of suitable surface treating methods include surface oxidation/reduction reaction, and coating by an organic compound or polymeric
25 compound. Skilled person in the art can easily prepare fiber embedded electrode structures according to this invention regardless of kinds of fiber materials or the process for pretreatment of the material. This means that the present invention is not limited to any specific types of fibers or the process for preparation of the fibriform.

Slurry for applying to an electrode is prepared by dispersing a conductive
30 fibriform (first conductive additive) and electrode active materials to a solvent, to which a proper organic binder, or binder dissolved in organic solvent is added. At

this time, carbon black other than fibriform or other kinds of carbonaceous powder (second conductive additive) can be mixed together in order to satisfy various objects considered in designing an electrode. The binder (of the present invention) includes polyvinylidene fluoride group, PVC group, PMMA group, SBR group, SBR Latex group, 5 PTFE (polytetrafluoroethylene) group, and rubber type polymer.

The organic solvent of the present invention is volatile and is not remained in the electrode after drying the electrode. Though nmp (n-methyl-2-pyrrolidone) or acetone and the like are used in general, but organic solvent is not limited in the present invention.

After the prepared slurry has been applied thinly on a current collector, the slurry 10 is heated in an oven to evaporate organic solvents. Thus, the electrode is completed. If necessary, the volume of an electrode could be decreased through compaction by roll press. Examples of a current collector include metal plate, metal foil, metal mesh, metal gauze, perforated metal foil, sintered metal fiber mesh, carbon paper, carbon sheet, carbon coated metal and the like. The present invention is not limited by the geometrical 15 structure, the elemental composition of the current collector.

The conductive fibriform, alone or along with a second conductor powder, can be dispersed in an organic solvent to facilitate dispersion of the slurry prior to preparing slurry mixing. Uniformly mixed slurry can be prepared by physical stirring or by ultrasonic dispersion.

20 The weight percentage of the conductive fibriform in the slurry can be from about 0.05% to about 50% of total weight of slurry, or from about 0.1% to about 50% of electrode total weight of active materials. In the case of using a secondary conductor powder, the rate of the powder weight to fibriform weight could change from about 0.01: 1 to about 1:0.01. However, preferable ratio of powder is within the range from about 25 0.1% to about 20% of the weight of active materials.

A cylindrical battery is provided using the electrode prepared as described above. In accordance with the known process, cathode and anode electrode terminals are welded and wound with a separator therebetween to prepare Jelly Roll. It is intercalated to a cylinder of 18650 scale. The electrode terminals are welded to the cylinder and its lid 30 and then an electrolyte is filled and sealed up so as to complete the battery.

DESCRIPTION OF THE PREFERRED

EMBODIMENT

The present invention will be illustrated in greater detail by way of the following examples. The examples are given for illustration of the invention and are not intended to be limiting thereof.

Example 1

5 PVdF was dissolved in nmp at a ratio of 9:1 to prepare an adhesive solution. LiCoO₂ (Seimi) and graphite (Lonza, KS6) were added to the solution, and mixed to prepare the slurry (Slurry A). The composition of the slurry A was LiCoO₂:graphite: PVdF=91:6:3. To the slurry A, a stainless steel fiber having a diameter of about 1.5 μ m and a length/diameter ratio of from about 100 to about 1000 was added, dispersed, and
10 mixed to prepare a slurry (Slurry B). The slurry B was applied on an aluminium foil and air-dried for about 30 minutes at 130 °C to dry the electrode.

The dispersion of metal fiber was examined by scanning electron microscopy and the surface photographs are shown in Figure 1.

On the contrary to a common disbelief that fibrous materials are difficult to
15 disperse within viscous media, the individual metal fibers were remarkably well distributed in the electrodes as shown in Figure 1.

Example 2

PVdF of 7.0g was dissolved to nmp of 65 ml, a synthesized graphite (MCMB) of 93.0g was added, stirred and dispersed (mixture A). a stainless steel (316L) fiber of 0.5g
20 having a diameter of 1.5 μ m and a length of about 0.5~1.0 mm was added to the mixture A and dispersed (mixture B). The composition of the mixture B less solvent was graphite: metal fiber: binder (PVdF)=91.2:2.0:6.8. After applying the mixture B on a copper foil at a thickness of 150 microns, it was first dried for 5 hours at an ambient temperature, dried again for 12 hours at about 130 °C, and then vacuum dried for 30
25 minutes at 130 °C. The micrographs of the electrode surface are also shown in figure 1.

Example 3

The electrode prepared as in example 1 was used as a positive electrode, each of lithium foils was used as a negative electrode and reference electrode, 1M LiClO₄/EC+DEC was used as an electrolyte solution. The cell was constructed. After
30 construction, voltage was stable as about 3.2 V

Discharging capacity according to each of cut-off voltage was shown in Table 1. The results of Fig. 2 and Table 1 illustrate that capacity of the electrode was considerably increased when a fibrous conductor was added. This result is because an efficient electron delivering passage was formed inside of the electrode. This allows a portion of electrode active materials once-isolated electrically to take part in cell reaction. Also, capacity according to cut-off voltage was shown in Table 1.

Table 1.

Changes of capacities of electrodes according to addition of a fibrous conductor

Content of Metal Fiber (%)	Capacities of Electrodes according to Cut-off Voltages (mAh/g)				
	4.0V	3.8V	3.6V	3.3V	3.0V
0	18	115	123	124	123
1	20 (11%)	124(8%)	130(6%)	131(6%)	132(6%)
3	24 (33%)	123(7%)	129(5%)	131(6%)	131(6%)
4	30 (67%)	128(11%)	132(7%)	133(7%)	134(7%)

(Values in parentheses are increasing ratios of capacities of electrodes)

This increasing of capacity means that the utility of active materials in the electrode is increased. Also, it means that active materials are not used to 100% before a fibrous metal conductor is added. It is understood that electron delivery by graphite particles as a conventional conductor is insufficient so that active materials partially isolated is electrically connected by metal fiber to take part in cell reaction. In the meantime, capacity is increased in proportion to the higher cut-off voltage of discharging curve. It means that IR drop of the electrode is decreased in proportion to the amount of metal added.

Example 4

The electrode prepared as in example 2 was used as a positive electrode, lithium foil was used as a negative electrode, and Celgard 1400 membrane was used as a separator. This layered structure and 1M LiPE6/EC+DEC as an electrolyte solution was used to make a cell. After completion of construction of the cell, voltage of an open circuit was about 3.2 V.

An intercalation/deintercalation of lithium ion was repeated using a current density per

unit area of 0.31 mA/cm^2 . The relationship between capacity and voltage of cell, and deintercalation capacity of carbon electrode are shown in Table 2, Table 3, Fig. 3.

Table 2.

Changes of high-speed discharging capacities according to addition of a fibrous conductor

(cut-off voltage = 1.5 V)

Types of Electrode	Capacities At Different Discharging Ratios (mAh/g carbon)				
	0.31	1.55	3.1	4.65	6.2mAh/cm ²
Electrode in which a fiber is added	300	274	229	265	139
Electrode in which a fiber is not added	281	257	230	149	98

Table 3.

Changes of high-speed discharging capacities according to addition of a fibrous conductor

(cut-off voltage = 0.9 V)

Types of Electrode	Capacities At Different Discharging Ratios (mAh/g carbon)				
	0.31	1.55	3.1	4.65	6.2mAh/cm ²
Electrode in which a fiber is added	297	272	224	174	128
Electrode in which a fiber is not added	279	256	226	126	73

The irreversible lithium ion uptake as well as reversible capacity from anodes have been examined and displayed in Figure 3. Compared to a control that contains no conductive additives, the fiber-containing electrode exhibited about 14% increase in reversible capacity, while the increase in irreversible reaction is only by 2 mAh/g. Carbon black-added anode with comparable volumetric quantity to fibers also

exhibits similar level of reversible capacity, but only at the expense of almost twofold increase in irreversible capacity toll.

The above results indicate that the contribution of metal fiber to the increase in the irreversible lithium uptake is almost none. The contribution from fiber to the electrode active area has been calculated and found negligible while the carbon black at the comparable volume content almost quadruples the electrode area. The increase in the reversible capacity is, again as in the case of cathodes, attributed to improved utilization of MCMB particles and lower IR drop that prevents the electrode from reaching to the cutoff potential prematurely.

Example 5

Metal fibrous conductor (a first conductor) was used along with a conventional powder type conductor (a second conductor). Nickel fiber of 0.4g having a diameter of $2\mu\text{m}$ and a length of 0.5 nm, carbon black of 0.4g, synthesized graphite (SFG15, Timcal) of 0.9g and solvent (nmp) of 20g were mixed and dispersed uniformly, and then mixed with 13 wt% PVdF/nmp solution of 10g and stirred by a pestle to give an uniform slurry.

The composition of the mixture excepts of the above described solvent was graphite: metal fiber: carbon black: binder (PVdF)=81:3.6:3.6:11.8. After applying the mixture on a copper foil at a thickness of 100 microns, the same processes as described in Example 2 were repeated to prepare the cell. It was conformed that the adhering strength of the electrode active materials was more increased. A high-speed discharging capacity was 8.3 mA/cm² which is 50% of a total capacity.

Example 6

After the mixture B prepared as in Example 1 was applied on Cu Exmet (mesh type collector) at a thickness of about 200 μm , the same processes as described in Example 2 were repeated to prepare the cell. It was conformed that the adhering strength of the electrode active materials was more increased. A high-speed discharging capacity was 8.3 mA/cm², which is 50% of a total capacity.

Example 7

Carbon whisker being undergone graphitization reaction of 0.5g having a diameter of $5\mu\text{m}$ and a length of 1 mm and a stainless steel fiber of 0.2g having a

diameter of 0.5 μ m and a length of 0.5 mm, MCMB of 9.0g and a solvent (nmp) of 17g were uniformly dispersed, mixed with 13wt% PVdF/nmp solution of 8.0g and stirred to prepare a slurry. After the mixture was applied on Cu foil at a thickness of about 100 microns, the drying process as described in Example 2 was performed to prepare the electrode for constructing a cell. The electrode prepared was appeared to have a very strong adhering strength. When recharging/discharging steps were repeated at 0.3 mA/cm², a capacity was maintained to about 95% of an initial capacity after 100 times of repetition.

Example 8

10 In order to verify the efficacy of fibrous conductive additives in real Li-ion battery applications, commercial-size cylindrical cells were manufactured using 316L fiber-added electrodes. The 18650 cells were compared at a nominal capacity of *ca.* 1400 mAh with all the electrode process variables and physical characteristics adjusted accordingly. The assembled batteries were subjected to cycle tests in the following order; i) 15 preconditioning at a 1/10C rate for the first charge and discharge sequence, ii) followed by four 1/5C rate cycles, and iii) subsequent 1C rate cycles with a restriction on the total charge time, which was 2.5 hours in the experiments shown in the Figure 5. During the 1C rate charge constant current mode with voltage limit of 4.1 V was applied.

20 For the first five cycles, where the charge/discharge current is low, the capacities for the two cells are similar. But when the current was raised to a 1C rate, it was evident that the capacity of the control cell sharply drops by more than 200 mAh and do not keep up with the metal fiber-added counterpart.

WHAT IS CLAIMED IS:

1. A rechargeable lithium ion battery comprising a cathode electrode, an anode electrode, an electrolyte and a separator, wherein at least one of said cathode electrode and anode electrode comprises an additive of from about 0.1 to about 50 % wt. of electrode active materials, and said additive is a fibriform having a diameter from about 0.1 to about 25 microns, and an aspect ratio of from about 4 to about 2500.
2. The battery according to Claim 1, wherein said additive comprises a material selected from the group consisting of iron, nickel, copper, zinc, titanium, aluminium, silver, gold, platinum, ion-chromium alloy, ion-chromium-nickel alloy, and aluminium alloy.
3. The battery according to Claim 1, wherein said additive comprises a fiber based on a synthesized graphite and an active carbon or a gas phase grown whisker type carbon fiber.
4. The battery according to Claim 1, wherein said cathode electrode comprises a material, as an active compound, selected from the group consisting of a compound capable of reacting reversibly with a lithium ion, a compound having a structure that a lithium ion can be intercalated, an organic sulfur compound, and a polymeric organic sulfur compound.
5. The battery according to Claim 4, wherein said active compound comprises active materials represented by:

$$\text{Li}_{1-x}\text{A}_x\text{Ni}_{1-y}\text{B}_y\text{O}_2$$
 (where, A= alkaline metal or alkaline earth metal,
 B=transition metal, $0 \leq x \leq 0.1$, $0 \leq y \leq 1.0$);
 $\text{LiMn}_{2-y}\text{M}_y\text{O}_4$ (where, M= Fe, Co, Ni: $0.02 \leq y \leq 0.3$);
 $\text{Li}_{1-x}\text{Ni}_{1-y}\text{B}_y\text{O}_2$ (where, B= transition metal. $0 \leq x \leq 0.1$, $0 \leq y \leq 1.0$);
 NbSe_3 ;
 $\text{Li}_x\text{V}_2\text{O}_5$; or
 $\text{Li}_x\text{V}_6\text{O}_{13}$ (where, $0 \leq x \leq 3.0$).
6. The battery according to Claim 1, wherein said anode

electrode comprises a material, as an active compound, selected from the group consisting of a compound capable of reacting reversibly with a lithium ion, a compound having a structure in which reaction products of a lithium ion with an electrolyte salt or an electrolyte solvent are capable of deposition or precipitation, a compound having a structure with which a lithium ion can be intercalated, lithium metal, lithium alloy, and carbon.

7. The battery according to Claim 1, wherein said additive comprises a metal fiber prepared from a material selected from the group of:

- a) a metal comprising iron, nickel, copper, zinc, titanium, aluminium, silver, gold, or platinum;
- b) ion-chromium-nickel alloy; and
- c) aluminium alloy including aluminium-copper, aluminium-manganese, aluminium-magnesium and aluminium-silicon-magnesium.

8. The battery according to Claim 1, wherein said electrolyte comprises a lithium salt selected from the group consisting of LiPF_6 , LiBF_4 , LiClO_4 , LiAsF_6 , LiSbF_6 , $\text{LiN}(\text{CF}_3\text{SO}_2)_2$, LiCF_3SO_2 , and $\text{LiN}(\text{SO}_2\text{C}_2\text{F}_5)_5$.

9. The battery according to Claim 1, wherein said electrolyte comprises a solvent selected from the group consisting of ethylene carbonate, propylene carbonate, vinylene carbonate, dimethyl carbonate, butylene carbonate, γ -butyrolactone, diethyl carbonate, ethylmethyl carbonate, N,N-dimethyl acetamide, dimethoxyethane and a mixture thereof.

10. The battery according to Claim 1, wherein said separator is a microporous polymeric membrane or nonfabric.

11. The battery according to Claim 1, wherein said metal fiber has a diameter from about 0.5 microns to about 4 microns, and an aspect ratio of from about 4 to about 2500.

12. The battery according to Claim 1, wherein said additive is a mixture of two or more different kind fibrous additives.

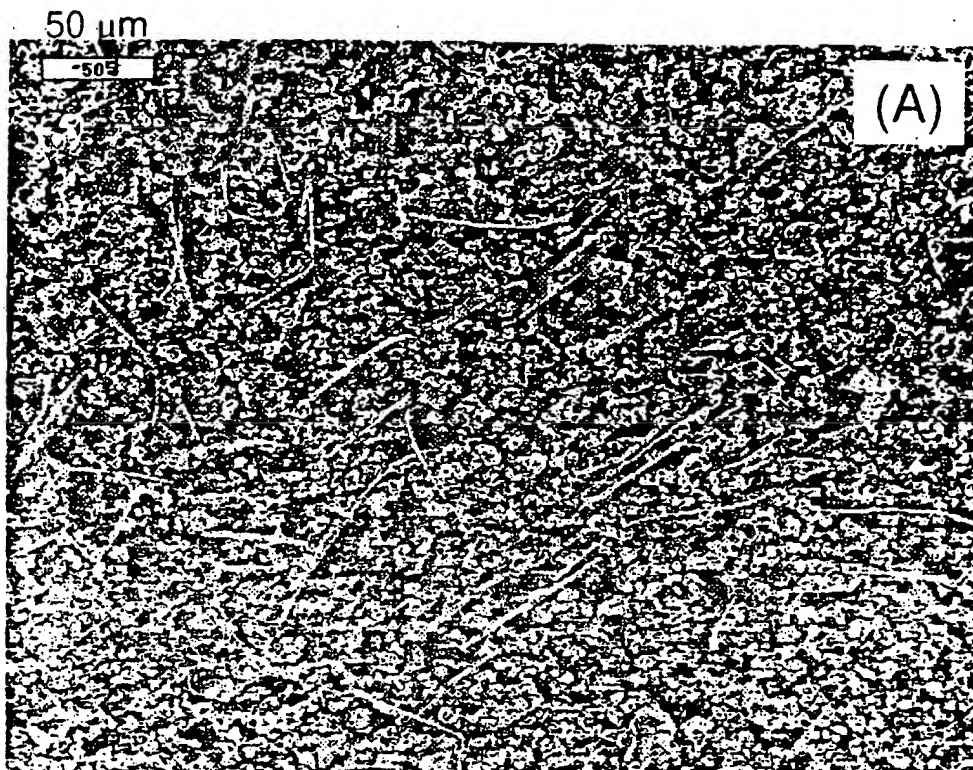
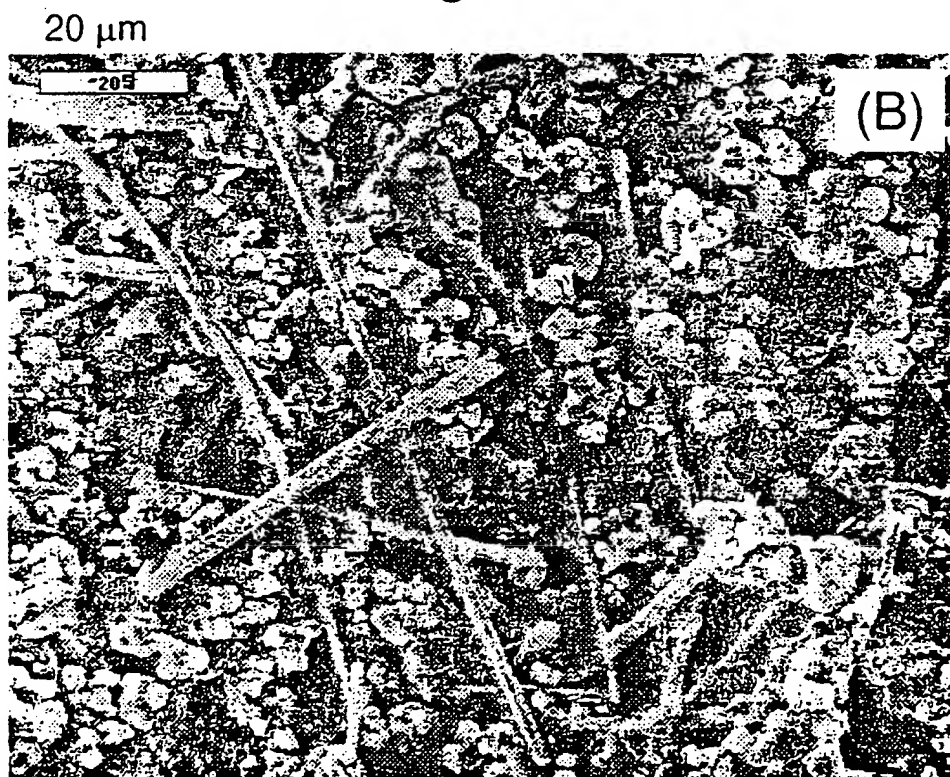
13. The battery according to Claim 1, wherein content of said additive is from about 0.1% to about 10% by weight of electrode active materials.

14. A method of manufacturing a rechargeable lithium

ion battery comprising the steps of:

- a) preparing a suspension by adding an additive of about 0.1% to about 50% by weight to electrode active materials, said additive consisting of a material selected from a group of iron, nickel, copper, zinc, titanium, aluminium, silver, gold,
5 platinum, ion-chromium alloy, ion-chromium-nickel alloy, and aluminium alloy and said additive being a fibriform having a diameter from about 0.5 to about 25 microns and an aspect ratio of from about 4 to about 2500;
- b) applying the suspension to a collector; and
- c) heating the collector obtained in step b).

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Fig. 1A ($\times 500$)Fig. 1B ($\times 1000$)

50 μ m

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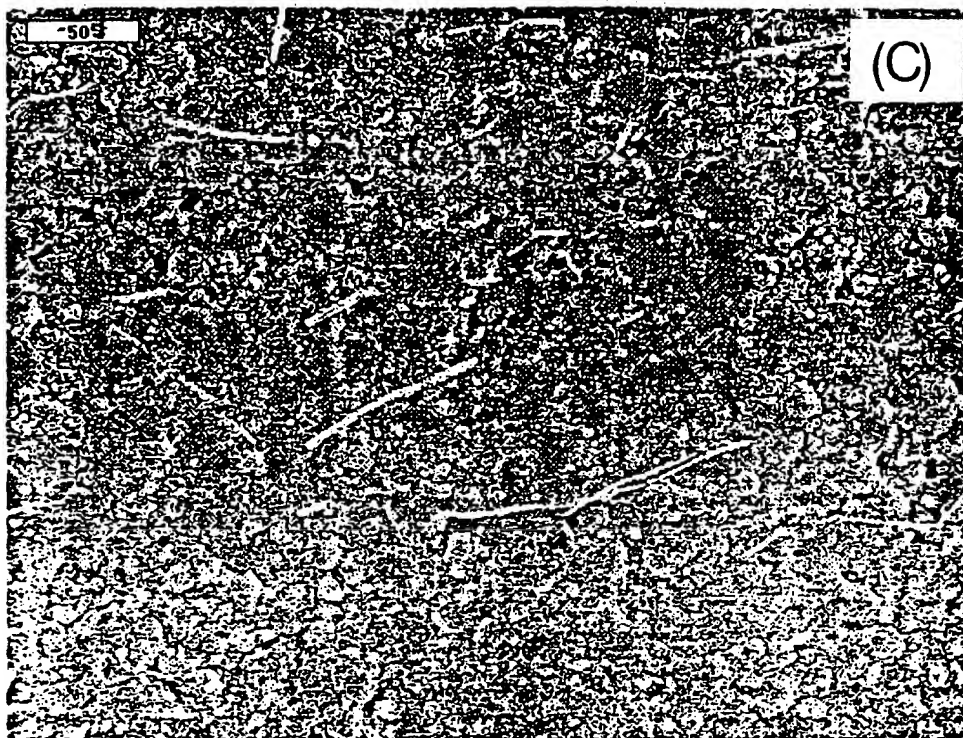


Fig. 1C

(X300)

5 μ m

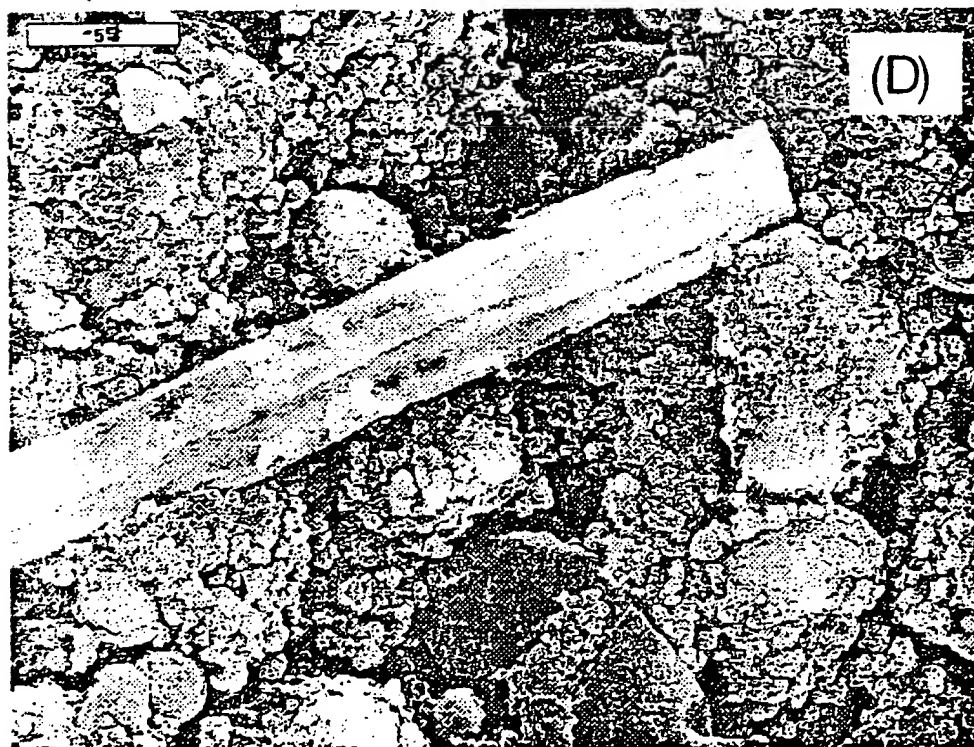
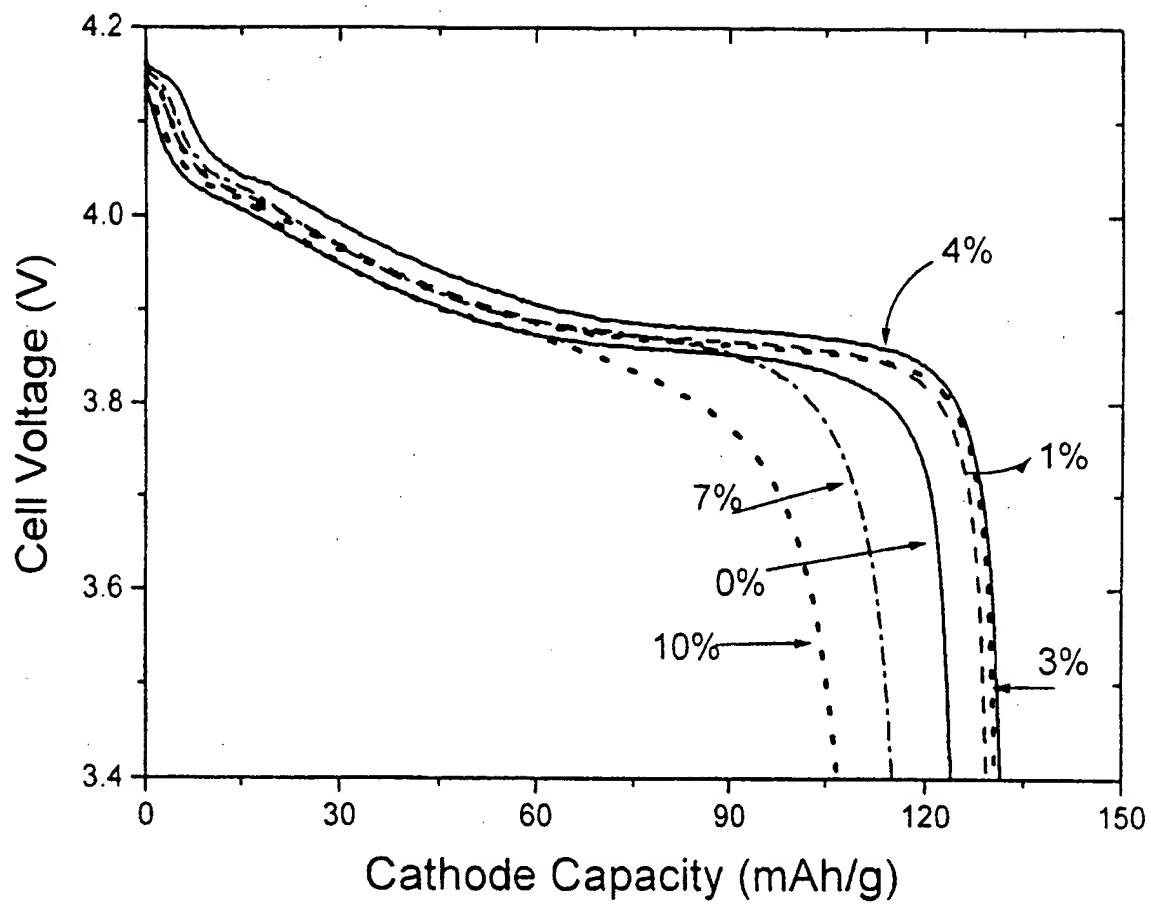


Fig. 1D

(x4000)

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F ig . 2

4/5

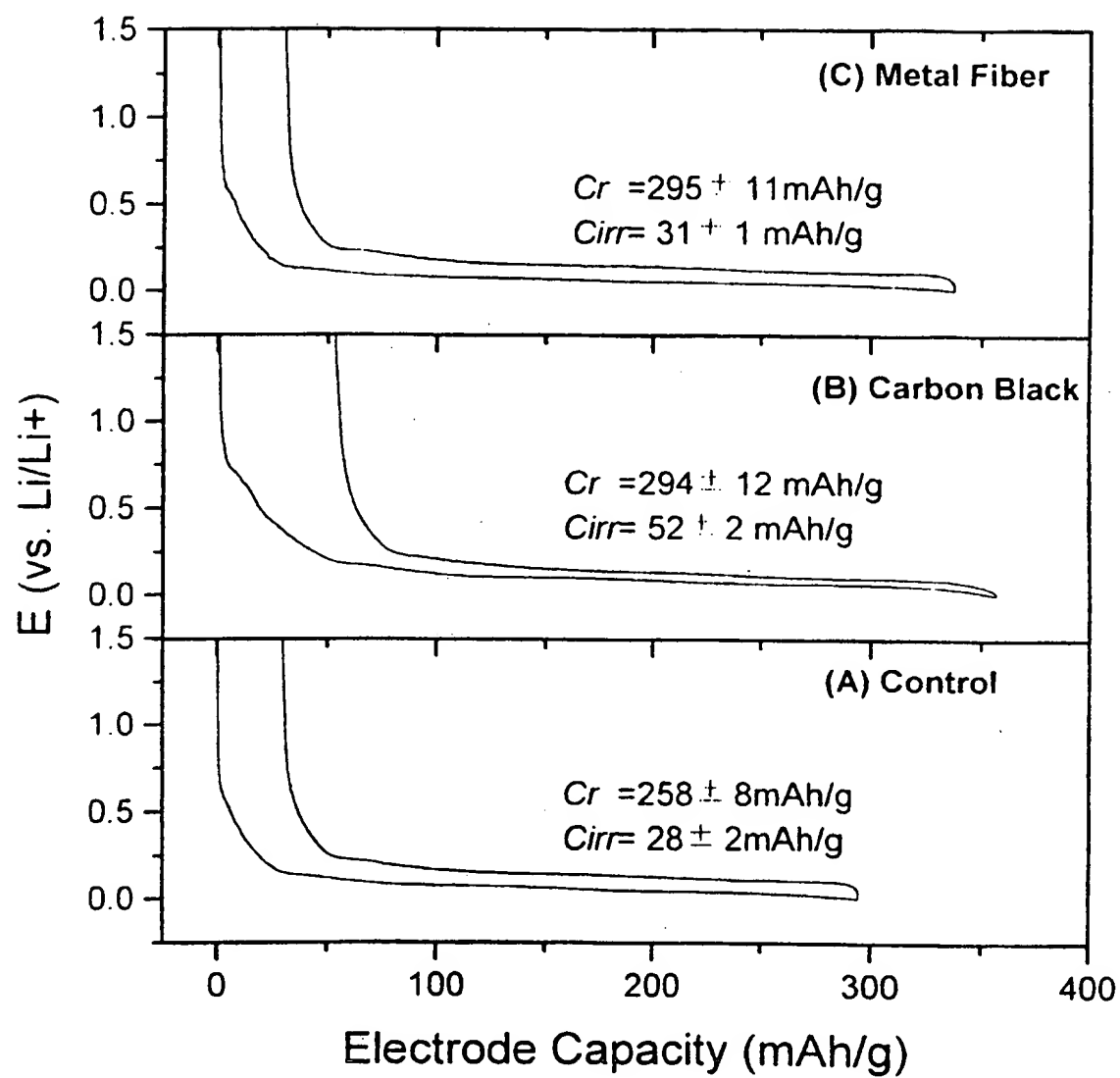


Fig. 3

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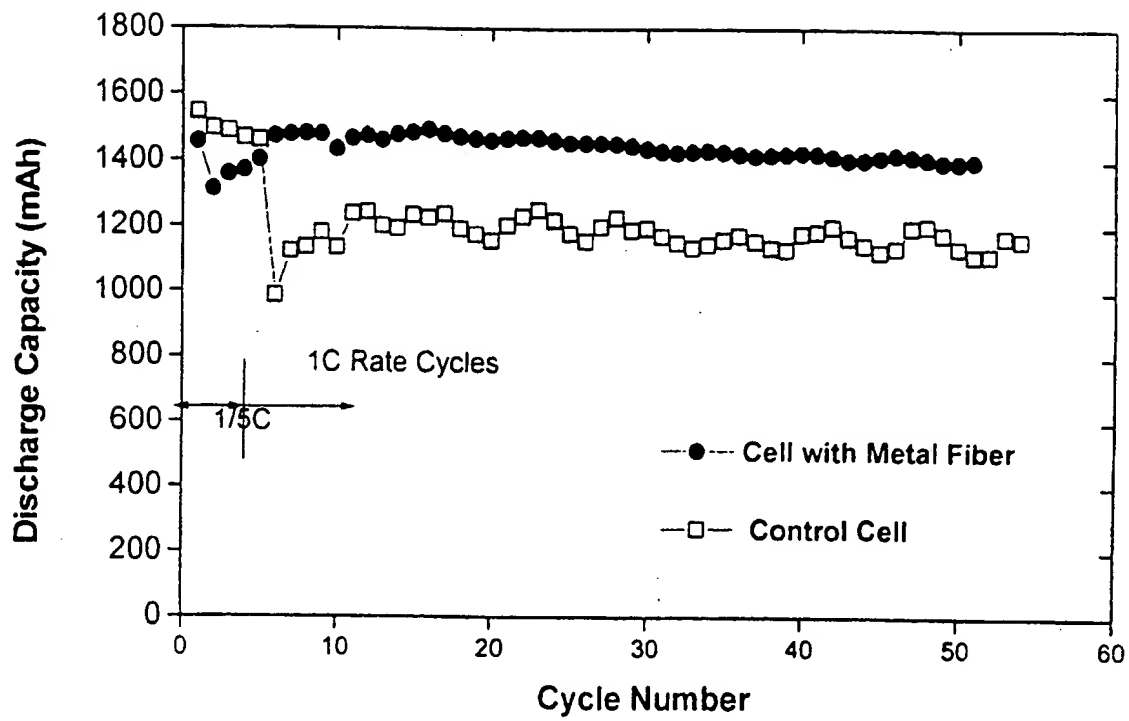


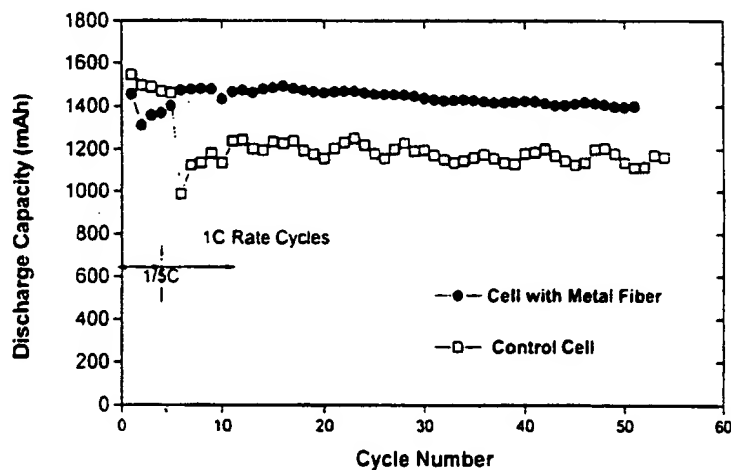
Fig. 4



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : H01M 10/40, 4/62		A3	(11) International Publication Number: WO 99/00001
			(43) International Publication Date: 7 January 1999 (07.01.99)
(21) International Application Number: PCT/KR98/00183		(74) Agent: KIM, Seong-Ki; Hubba Hubba Building, 7th floor, 648, Yoksam-dong, Kangnam-gu, Seoul 135-080 (KR).	
(22) International Filing Date: 26 June 1998 (26.06.98)		(81) Designated States: CN, JP, US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).	
(30) Priority Data: 1997/28473 27 June 1997 (27.06.97) KR 1997/28474 27 June 1997 (27.06.97) KR 1998/2226 24 January 1998 (24.01.98) KR		Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(71) Applicant (for all designated States except US): LG CHEMICAL, LTD. [KR/KR]; LG Twin Towers, 20, Yoido-dong, Yongsongpo-gu, Seoul 150-721 (KR).		(88) Date of publication of the international search report: 30 September 1999 (30.09.99)	
(72) Inventors; and (75) Inventors/Applicants (for US only): AHN, Soon-Ho [KR/KR]; 107-1106, Lucky Hana Apt., Shinsung-dong, Yousung-gu, Taejon 305-345 (KR). CHUNG, Geun-Chang [KR/KR]; 6-103, LG Apt., Doryong-dong, Yousung-gu, Taejon 305-345 (KR). LIM, Mi-Ra [KR/KR]; 3-103, LG Chem. Dormitory, Doryong-dong, Yousung-gu, Taejon 305-340 (KR). JUN, Sung-Hui [KR/KR]; 3-103, LG Chem. Dormitory, Doryong-dong, Yousung-gu, Taejon 305-340 (KR). BAE, Joon-Sung [KR/KR]; 3-412, LG Chem. Dormitory, Doryong-dong, Yousung-gu, Taejon 305-340 (KR). KWAK, Mi-Seon [KR/KR]; 105-1110, Hanbat Garden, Sanjung-dong, Jung-gu, Taejon 301-211 (KR).			

(54) Title: LITHIUM ION SECONDARY BATTERY AND MANUFACTURING METHOD OF THE SAME



(57) Abstract

The present invention discloses an improved lithium battery comprising a positive electrode, a negative electrode and a separator, and a manufacturing method of the same. The positive and negative electrodes comprise an active compounds selected from the group consisting of a compound being capable of reacting reversibly with a lithium ion, a compound having a structure that products of reacting a lithium ion with an electrolyte salt or an electrolyte solvent are capable of deposition or precipitation, or a compound having a structure that a lithium ion can be intercalated, an additive comprising a fiber of micron scale having an electron conductivity, a collector comprising metal or carbon material, and an adhesive. The lithium ion secondary battery prepared from the above described electrode has a low internal resistance, a high capacity and a high-speed recharging-discharging capability.

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/KR 98/00183

A. CLASSIFICATION OF SUBJECT MATTER

IPC⁶: H 01 M 10/40, 4/62

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C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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A	EP 0 717 457 A2 (KATAYAMA SPECIAL INDUSTRIES, LTD), 19 June 1996 (19.06.96), claims 1,15.	1-14
A	Database WPI on EPO, week 9531, London: Derwent Publications Ltd., AN 95-234877, class W 04 X 16, JP 7-142 052 A (SUMITOMO ELECTRIC IND CO), abstract.	1-14
A	Patent Abstracts of Japan, Vol.12, No.357 (E-662), 1988, JP 63-114 057 A (SANYO ELECTRIC CO LTD), abstract.	1-14

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JP A2 9161807	20-06-1997	keine - none - rien	
EP A2 717457	19-06-1996	CA AA 2163819	29-05-1996
		CA AA 2254504	29-05-1996
		CA AA 2254551	29-05-1996
		CA AA 2254640	29-05-1996
		CN A 1127433	24-07-1996
		EP A3 717457	08-01-1997
		JP A2 8213026	20-08-1996
JP A2 7142052	02-06-1995	keine - none - rien	
JP A2 63114057	18-05-1988	keine - none - rien	